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## Amendments to Claims

(Currently Amended): A method for controlling arthropods comprising contacting the arthropods or their environment with an arthropodicidally effective amount of a compound of Formula 1, its N-oxide or agriculturally suitable salts

wherein

A and B are independently O or S;

each J-is independently a phenyl or naphthyl group substituted with 1 to 2 R5 and optionally substituted with 1 to 3 R6;

or each J is independently a 5 or 6-membered heteroaromatic ring or an aromatic 8-, 9or 10-membered fused heterobicyclic ring system-wherein each ring or ring system is optionally substituted with 1 to 4 R7;

n is 1 to 4;

 $R^1$  is H; or  $C_1$ - $C_6$  alkyl,  $C_2$ - $C_6$  alkenyl,  $C_2$ - $C_6$  alkynyl or  $C_3$ - $C_6$  cycloalkyl each optionally substituted with one or more substituents selected from the group consisting of halogen, CN, NO2, hydroxy, C1-C4 alkoxy, C1-C4 alkylthio, C1-C4 alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>2</sub>-C<sub>4</sub> alkoxycarbonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino and C<sub>3</sub>-C<sub>6</sub> cycloalkylamino; or

R1 is C2-C6 alkylcarbonyl, C2-C6 alkoxycarbonyl, C2-C6 alkylaminocarbonyl, C3-C8 dialkylaminocarbonyl or C(=A)J;

 $R^2$  is H,  $C_1$ - $C_6$  alkyl,  $C_2$ - $C_6$  alkenyl,  $C_2$ - $C_6$  alkynyl,  $C_3$ - $C_6$  cycloalkyl,  $C_1$ - $C_4$  alkoxy, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl or C2-C6 alkylcarbonyl;

 $R^3$  is H; G;  $C_1$ - $C_6$  alkyl,  $C_2$ - $C_6$  alkenyl,  $C_2$ - $C_6$  alkynyl,  $C_3$ - $C_6$  cycloalkyl, each optionally substituted with one or more substituents selected from the group consisting of halogen, G, CN, NO2, hydroxy, C1-C4 alkoxy, C1-C4 haloalkoxy,  $C_1-C_4 \text{ alkylthio, } C_1-C_4 \text{ alkylsulfinyl, } C_1-C_4 \text{ alkylsulfonyl, } C_2-C_6 \text{ alkoxycarbonyl, }$ C2-C6 alkylcarbonyl, C3-C6 trialkylsilyl, and a phenyl, phenoxy or 5- or 6membered heteroaromatic ring, each ring optionally substituted with one to three

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substituents independently selected from the group consisting of C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>2</sub>-C<sub>4</sub> haloalkenyl, C<sub>2</sub>-C<sub>4</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, C<sub>3</sub>-C<sub>6</sub> (alkyl)cycloalkylamino, C<sub>2</sub>-C<sub>4</sub> alkylcarbonyl, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub> dialkylaminocarbonyl and C<sub>3</sub>-C<sub>6</sub> trialkylsilyl; C<sub>1</sub>-C<sub>4</sub> alkoxy; C<sub>1</sub>-C<sub>4</sub> alkylamino; C<sub>2</sub>-C<sub>8</sub> dialkylamino; C<sub>3</sub>-C<sub>6</sub> cycloalkylamino; C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl or C<sub>2</sub>-C<sub>6</sub> alkylcarbonyl; or

- R<sup>2</sup> and R<sup>3</sup> can be taken together with the nitrogen to which they are attached to form a ring containing 2 to 6 atoms of carbon and optionally one additional atom of nitrogen, sulfur or oxygen, said ring may be optionally substituted with 1 to 4 substituents selected from the group consisting of C<sub>1</sub>-C<sub>2</sub> alkyl, halogen, CN, NO<sub>2</sub> and C<sub>1</sub>-C<sub>2</sub> alkoxy;
- G is a 5- or 6-membered nonaromatic carbocyclic or heterocyclic ring, optionally including one or two ring members selected from the group consisting of C(=O), SO or S(O)<sub>2</sub> and optionally substituted with 1 to 4 substituents selected from the group consisting of C<sub>1</sub>-C<sub>2</sub> alkyl, halogen, CN, NO<sub>2</sub> and C<sub>1</sub>-C<sub>2</sub> alkoxy;
- each R<sup>4</sup> is independently H, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>6</sub> haloalkyl, C<sub>2</sub>-C<sub>6</sub> haloalkenyl, C<sub>2</sub>-C<sub>6</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO<sub>2</sub>, hydroxy, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> haloalkylthio, C<sub>1</sub>-C<sub>4</sub> haloalkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> haloalkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, or C<sub>3</sub>-C<sub>6</sub> trialkylsilyl; or
- each R<sup>4</sup> is independently phenyl, benzyl or phenoxy, each optionally substituted with C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>2</sub>-C<sub>4</sub> haloalkenyl, C<sub>2</sub>-C<sub>4</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, C<sub>3</sub>-C<sub>6</sub> (alkyl)cycloalkylamino, C<sub>2</sub>-C<sub>4</sub> alkylcarbonyl, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub> dialkylaminocarbonyl or C<sub>3</sub>-C<sub>6</sub> trialkylsilyl;
- each  $R^5$  is independently  $C_1$   $C_6$  alkyl,  $C_2$   $C_6$  alkenyl,  $C_2$   $C_6$  alkynyl,  $C_3$   $C_6$  cycloalkyl,  $C_1$   $C_6$  haloalkyl,  $C_2$   $C_6$  haloalkenyl,  $C_2$   $C_6$  haloalkynyl,  $C_3$   $C_6$  haloalkyl, halogen, CN,  $CO_2H$ ,  $CONH_2$ ,  $NO_2$ , hydroxy,  $C_1$   $C_6$  alkoxy,  $C_1$   $C_6$  haloalkoxy,  $C_1$   $C_6$  alkylthio,  $C_1$   $C_6$  alkylsulfinyl,  $C_1$   $C_6$  alkylsulfonyl,  $C_1$   $C_6$  haloalkylsulfinyl,  $C_1$   $C_6$  haloalkylsulfonyl,  $C_1$   $C_6$  alkylamino,  $C_2$   $C_{12}$  dialkylamino,  $C_3$   $C_6$  cycloalkylamino,  $C_4$   $C_6$  alkylamino,  $C_5$   $C_6$  alkylamino,  $C_6$   $C_6$  C

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alkoxycarbonyl, C2 C6 alkylaminocarbonyl, G2-C8 dialkylaminocarbonyl, or C3-C6 triallty sily 1; or

- (R5)2 when attached to adjacent earbon atoms can be taken together as OCF2O, -CF<sub>2</sub>CF<sub>2</sub>O , or OCF<sub>2</sub>CF<sub>2</sub>O ;
- each R6 is independently H, halogen, C1-C6 alkyl, C2-C6 alkenyl, C2-C6-alkynyl, C2-C6 eyeloalkyl, C1 G4 alkoxy or C2 C4 alkoxycarbonyl; or
- each R6 is independently a phenyl, benzyl, phenoxy, 5 or 6 membered heteroaromatic ring or an aromatic-8, 9- or 10 membered fused heterobioyelie ring-system, each ring optionally substituted with one to three substituents independently selected from the group consisting of C1 C4 alkyl, C2 C4 alkenyl, C2 C4 alkynyl, C3 C6 eyelealkyl, C<sub>1</sub>-C<sub>4</sub> halealkyl, C<sub>2</sub>-C<sub>4</sub>-halealkenyl, C<sub>2</sub>-C<sub>4</sub> halealkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO2, C1-C4 alkoxy, C1-C4 haloalkoxy, C1-C4 alkylthio, C1-C4-alkylsulfinyl, C1-C4-alkylsulfonyl, C1-C4 alkylamino, C2-C8  $\frac{\text{dialkylamino, } C_3 \cdot C_6 \cdot \text{cycloalkylamino, } C_2 \cdot C_6 \cdot \text{(alkyl)cycloalkylamino, } C_2 \cdot C_4}{\text{dialkylamino, } C_3 \cdot C_6 \cdot \text{(alkyl)cycloalkylamino, } C_2 \cdot C_6}$ alkylearbonyl, C2 C6 alkoxyoarbonyl, C2 C6 alkylaminocarbonyl, C3 C8 dialkylaminecarbonyl and C2 C6 trialkylsilyl;
- each R7 is independently H, C1-C6 alkyl, C2-C6 alkenyl, C2-C6 alkynyl, C3-C6 cycloalkyl,  $C_1$ - $C_6$  haloalkyl,  $C_2$ - $C_6$  haloalkenyl,  $C_2$ - $C_6$  haloalkynyl,  $C_3$ - $C_6$ halocycloalkyl, halogen, CN, CO<sub>2</sub>H, CONH<sub>2</sub>, NO<sub>2</sub>, hydroxy, C<sub>1</sub>-C<sub>4</sub> alkoxy,  $C_1$ - $C_4$  haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$  haloalkylthio,  $C_1$ - $C_4$  haloalkylsulfinyl,  $C_1$ - $C_4$  haloalkylsulfonyl,  $C_1$ - $C_4$ alkylamino, C2-C8 dialkylamino, C3-C6 cycloalkylamino, C2-C6 alkylcarbonyl, C2-C6 alkoxycarbonyl, C2-C6 alkylaminocarbonyl, C3-C8 dialkylaminocarbonyl, or C3-C6 trialkylsilyl; or
- each R7 is independently a phenyl, benzyl, benzoyl, phenoxy, 5- or 6-membered heteroaromatic ring or an aromatic 8-, 9- or 10-membered fused heterobicyclic ring system, each ring optionally substituted with one to three substituents independently selected from the group consisting of C1-C4 alkyl, C2-C4 alkenyl,  $C_2$ - $C_4$  alkynyl,  $C_3$ - $C_6$  cycloalkyl,  $C_1$ - $C_4$  haloalkyl,  $C_2$ - $C_4$  haloalkenyl,  $C_2$ - $C_4$ haloalkynyl,  $C_3$ - $C_6$  halocycloalkyl, halogen, CN,  $NO_2$ ,  $C_1$ - $C_4$  alkoxy,  $C_1$ - $C_4$ haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$ alkylamino, C2-C8 dialkylamino, C3-C6 cycloalkylamino, C3-C6 (alkyl)cycloalkylamino, C2-C4 alkylcarbonyl, C2-C6 alkoxycarbonyl, C2-C6 alkylaminocarbonyl, C3-C8 dialkylaminocarbonyl and C3-C6 trialkylsilyl;

provided that

(1) when A and B-are both O, R2 is H or C1 - C3 alkyl, R3 is H or C1 - C3 alkyl and R4 is II, halogon, C1-C6 alkyl, phenyl, hydroxy or C1-C6 alkoxy, then one R5 is other than halogen, C1-C6 alkyl, hydroxy or C1-C6 alkoxy; or

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- (2) Jis other than an optionally 1,2,3 thindiazole.
- (3) when J is an optionally substituted 5 membered heteroaromatic ring, then R<sup>2</sup> and R<sup>2</sup> are taken together with the nitrogen to which they are attached to form a ring containing 2 to 6 atoms of carbon and optionally one additional atom of nitrogen, sulfur or oxygen, said ring may be optionally substituted with 1 to 4 substituents selected from the group consisting of C<sub>1</sub> C<sub>2</sub> alkyl, halogen, CN, NO<sub>2</sub> and C<sub>1</sub>-C<sub>2</sub> alkoxy.
- 2. (Withdrawn): The method of Claim 1 wherein J is a phenyl group substituted with 1 to 2 R<sup>5</sup> and optionally substituted with 1 to 3 R<sup>6</sup>.
  - 3. (Withdrawn): The method of Claim 2 wherein

A and B are both O;

n is 1 to 2;

- $R^1$  is H,  $C_1$ - $C_4$  alkyl,  $C_2$ - $C_4$  alkenyl,  $C_2$ - $C_4$  alkynyl,  $C_3$ - $C_6$  cycloalkyl,  $C_2$ - $C_6$  alkylcarbonyl or  $C_2$ - $C_6$  alkoxycarbonyl;
- R<sup>2</sup> is H, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>2</sub>-C<sub>6</sub> alkylcarbonyl or C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl;
- R<sup>3</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl or C<sub>3</sub>-C<sub>6</sub> cycloalkyl each optionally substituted with one or more substituents selected from the group consisting of halogen, CN, C<sub>1</sub>-C<sub>2</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub> alkylthio, C<sub>1</sub>-C<sub>2</sub> alkylsulfinyl and C<sub>1</sub>-C<sub>2</sub> alkylsulfonyl;
- one of the R<sup>4</sup> groups is attached to the phenyl ring at the 2-position or 5-position, and said R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> haloalkylsulfinyl, or C<sub>1</sub>-C<sub>4</sub> haloalkylsulfonyl;
- each  $R^5$  is independently  $C_1$ - $C_4$  haloalkyl, CN, NO<sub>2</sub>,  $C_1$ - $C_4$  haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$  haloalkylsulfonyl,  $C_1$ - $C_4$  haloalkylsulfonyl or  $C_2$ - $C_4$  alkoxycarbonyl; or
- $(R^5)_2$  when attached to adjacent carbon atoms can be taken together as -OCF $_2$ O-, -CF $_2$ CF $_2$ O- or -OCF $_2$ CF $_2$ O-; and
- each R6 is independently H, halogen, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>2</sub> alkoxy or C<sub>2</sub>-C<sub>4</sub> alkoxycarbonyl, or
- each R6 is independently a phenyl or a 5- or 6-membered heteroaromatic ring, each ring optionally substituted with C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>2</sub>-C<sub>4</sub> haloalkenyl, C<sub>2</sub>-C<sub>4</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, C<sub>3</sub>-C<sub>6</sub> (alkyl)cycloalkylamino, C<sub>2</sub>-C<sub>4</sub>

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alkylcarbonyl,  $C_2$ - $C_6$  alkoxycarbonyl,  $C_2$ - $C_6$  alkylaminocarbonyl,  $C_3$ - $C_8$ dialkylaminocarbonyl or C3-C6 trialkylsilyl.

(Withdrawn): The method of Claim 3 wherein

R1 and R2 are both H;

 $R^3$  is  $C_1$ - $C_4$  alkyl optionally substituted with halogen, CN, OCH<sub>3</sub>, or  $S(O)_pCH_3$ ; each R4 is independently H, CH3, CF3, OCF3, OCHF2, S(O)pCF3, S(O)pCHF2, CN or halogen;

each R<sup>5</sup> is independently CF<sub>3</sub>, OCF<sub>3</sub>, OCHF<sub>2</sub>, S(O)<sub>p</sub>CF<sub>3</sub>, S(O)<sub>p</sub>CHF<sub>2</sub>, OCH<sub>2</sub>CF<sub>3</sub>, OCF<sub>2</sub>CHF<sub>2</sub>, S(O)<sub>p</sub>CH<sub>2</sub>CF<sub>3</sub> or S(O)<sub>p</sub>CF<sub>2</sub>CHF<sub>2</sub>;

each R6 is independently H, halogen or methyl; or phenyl, pyrazole, imidazole, triazole, pyridine or pyrimidine, each ring optionally substituted with C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, halogen or CN; and

p is 0, 1 or 2.

- (Withdrawn): The method of Claim 4 wherein  $\mathbb{R}^3$  is *i*-propyl or *t*-butyl. 5.
- (Canceled). 6.
- (Currently Amended): The method of Claim 6 1 wherein 7.

J is a 5- or 6-membered heteroaromatic ring selected from the group consisting of J-1, J-2, J-3, J-4 and J-5, wherein J-1 and J-2-are optionally substituted with 1-to 3 R<sup>7</sup> and J-3, J 4 and J-5 are substituted with R7

Q is O, S or NR7; and

W, X, Y and Z are independently N or CR7, provided that in-J-4 and J-5 at least one of W. X. Y or Z is N.

(Currently Amended): The method of Claim 6 or 7 wherein

A and B are O;

n is 1 to 2;

 $R^1$  is H,  $C_1$ - $C_4$  alkyl,  $C_2$ - $C_4$  alkenyl,  $C_2$ - $C_4$  alkynyl,  $C_2$ - $C_6$  alkylcarbonyl or  $C_2$ - $C_6$ aikoxycarbonyl;

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- R2 is H, C1-C4 alkyl, C2-C4 alkenyl, C2-C4 alkynyl, C3-C6 cycloalkyl, C2-C6 alkylcarbonyl or C2-C6 alkoxycarbonyl;
- R3 is H; or C1-C6 alkyl, C2-C6 alkenyl, C2-C6 alkynyl or C3-C6 cycloalkyl each optionally substituted with one or more substituents selected from the group consisting of halogen, CN, C1-C2 alkoxy, C1-C2 alkylthio, C1-C2 alkylsulfinyl and  $C_1$ - $C_2$  alkylsulfonyl;
- one of the R<sup>4</sup> groups is attached to the phenyl ring at the 2-position, and said R<sup>4</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$  haloalkylthio,  $C_1$ - $C_4$  haloalkylsulfinyl or  $C_1$ - $C_4$  haloalkylsulfonyl; and
- each R7 is independently H, C1-C4 alkyl, C1-C4 haloalkyl, halogen, CN, NO2, C1-C4 haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$ haloalkylthio, C1-C4 haloalkylsulfinyl, C1-C4 haloalkylsulfonyl or C2-C4 alkoxycarbonyl; or a phenyl or a 5- or 6-membered heteroaromatic ring, each ring optionally substituted with C1-C4 alkyl, C2-C4 alkenyl, C2-C4 alkynyl,  $C_3$ - $C_6$  cycloalkyl,  $C_1$ - $C_4$  haloalkyl,  $C_2$ - $C_4$  haloalkenyl,  $C_2$ - $C_4$  haloalkynyl,  $C_3$ - $C_6$ halocycloalkyl, halogen, CN, NO2, C1-C4 alkoxy, C1-C4 haloalkoxy, C1-C4 alkylthio, C1-C4 alkylsulfinyl, C1-C4 alkylsulfonyl, C1-C4 alkylamino, C2-C8 dialkylamino, C3-C6 cycloalkylamino, C3-C6 (alkyl)cycloalkylamino, C2-C4 alkylcarbonyl, C2-C6 alkoxycarbonyl, C2-C6 alkylaminocarbonyl, C3-C8 dialkylaminocarbonyl or C3-C6 trialkylsilyl.
  - (Currently Amended): The method of Claim 8 wherein J is selected from the 9. group consisting of pyridine, and pyrimidine, pyrazole, imidazole, triazoles, thiophene and thiazole, each optionally substituted with 1 to 3 R7.
- (Currently Amended): The method of Claim 9 wherein
- J is selected from the group consisting of pyridine, and pyrimidine, pyrazole, thiophene and thiazole, each optionally substituted with 1 to 3 R7;

R1 and R2 are both H;

- R<sup>3</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with halogen, CN, OCH<sub>3</sub>, or S(O)<sub>p</sub>CH<sub>3</sub>; each R4 is independently H, CH3, CF3, OCF3, OCHF2, S(O)pCF3, S(O)pCHF2, CN or halogen;
- each R<sup>7</sup> is independently H, halogen, CH<sub>3</sub>, CF<sub>3</sub>, OCHF<sub>2</sub>, S(O)<sub>p</sub>CF<sub>3</sub>, S(O)<sub>p</sub>CHF<sub>2</sub>, OCH<sub>2</sub>CF<sub>3</sub>, OCF<sub>2</sub>CHF<sub>2</sub>, S(O)<sub>p</sub>CH<sub>2</sub>CF<sub>3</sub>, S(O)<sub>p</sub>CF<sub>2</sub>CHF<sub>2</sub>; or phenyl, pyrazole, imidazole, triazole, pyridine or pyrimidine, each ring optionally substituted with  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_4$  haloalkyl,  $C_1$ - $C_4$  alkoxy,  $C_1$ - $C_4$  haloalkoxy,  $C_1$ - $C_4$  alkylthio, C1-C4 alkylsulfinyl, C1-C4 alkylsulfonyl, halogen or CN; and p is 0, 1 or 2.

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- 11. (Original): The method of Claim 10 wherein J is a pyridine optionally substituted with 1 to 3 R<sup>7</sup>.
- 12. (Original): The method of Claim 11 wherein one  $\mathbb{R}^7$  is a phenyl optionally substituted with  $\mathbb{C}_1$ - $\mathbb{C}_4$  alkyl,  $\mathbb{C}_1$ - $\mathbb{C}_4$  haloalkyl, halogen or  $\mathbb{C}\mathbb{N}$ .
- 13. (Original): The method of Claim 11 wherein one  $\mathbb{R}^7$  is a pyrazole, imidazole, triazole, pyridine or pyrimidine, each ring optionally substituted with  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_4$  haloalkyl, halogen or CN.
- 14. (Original): The method of Claim 10 wherein J is a pyrimidine optionally substituted with 1 to 3 R<sup>7</sup>.
- 15. (Original): The method of Claim 14 wherein one  $\mathbb{R}^7$  is a phenyl optionally substituted with  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_4$  haloalkyl, halogen or CN.
- 16. (Original): The method of Claim 14 wherein one  $\mathbb{R}^7$  is a pyrazole, imidazole, triazole, pyridine or pyrimidine, each ring optionally substituted with  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_4$  haloalkyl, halogen or CN.
  - 17. (Canceled).
  - 18. (Canceled).
  - 19. (Canceled).
  - 20. (Canceled).
- 21. (Currently Amended): The method of Claim 1 comprising a compound of Formula 1 selected from the group consisting of: which is

3 methyl N (1 methylethyl) 2 [[4 (trifluoremethyl)benzoyl]amino] benzamide,

2-methyl N-[2-methyl-6-[[(1-methylethyl)amino]earbonyl]phenyl] 4-

(trifluoromethyl)benzamide, and

2-methyl-N-[2-methyl-6-[[(1-methylethyl)amino]carbonyl]phenyl]-6-(trifluoromethyl)-3-pyridinecarboxamide.

22. (Currently Amended): A compound of Formula 1, its N-oxides and agriculturally suitable salts

wherein

A and B are independently O or S;

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- each-J is independently a phenyl or naphthyl group substituted with 1 to 2 R<sup>5</sup> and eptionally substituted with 1 to 3 R<sup>6</sup>;
- or each J is independently a 5 or 6-membered heteroaromatic ring or an aromatic 8,9or 10 membered fused heterobic ring system wherein each ring or ring system is optionally substituted with 1 to 4 R<sup>7</sup>;

n is 1 to 4;

- R<sup>1</sup> is H; or C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl or C<sub>3</sub>-C<sub>6</sub> cycloalkyl each optionally substituted with one or more substituents selected from the group consisting of halogen, CN, NO<sub>2</sub>, hydroxy, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>2</sub>-C<sub>4</sub> alkoxycarbonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino and C<sub>3</sub>-C<sub>6</sub> cycloalkylamino; or
- R1 is C<sub>2</sub>-C<sub>6</sub> alkylcarbonyl, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub> dialkylaminocarbonyl or C(=A)J;
- R<sup>2</sup> is H, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl or C<sub>2</sub>-C<sub>6</sub> alkylcarbonyl;
- R³ is H; C₁-C6 alkyl, C₂-C6 alkenyl, C₂-C6 alkynyl, C₃-C6 cycloalkyl, each optionally substituted with one or more substituents selected from the group consisting of halogen, CN, NO₂, hydroxy, C₁-C₄ alkoxy, C₁-C₄ haloalkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, C₂-C6 alkoxycarbonyl, C₂-C6 alkylcarbonyl, C₃-C6 trialkylsilyl, and a phenoxy ring optionally substituted with one to three substituents independently selected from the group consisting of C₁-C₄ alkyl, C₂-C₄ alkenyl, C₂-C₄ alkynyl, C₃-C6 cycloalkyl, C₁-C₄ haloalkyl, C₂-C₄ haloalkenyl, C₂-C₄ haloalkynyl, C₃-C6 halocycloalkyl, halogen, CN, NO₂, C₁-C₄ alkoxy, C₁-C₄ haloalkoxy, C₁-C₄ alkylthio, C₁-C₄ alkylsulfinyl, C₁-C₄ alkylsulfonyl, C₁-C₄ alkylamino, C₂-Cβ dialkylamino, C₃-C6 cycloalkylamino, C₃-C6 (alkyl)cycloalkylamino, C₂-Cβ dialkylamino, C₂-C6 alkoxycarbonyl, C₂-C6 alkylaminocarbonyl, C₃-Cβ dialkylamino; C₃-C6 cycloalkylamino; C₂-Cβ alkoxycarbonyl or C₂-Cβ dialkylamino; C₃-C6 cycloalkylamino; C₂-Cβ alkoxycarbonyl or C₂-Cβ alkylamino; C₃-Cβ cycloalkylamino; C₂-Cβ alkoxycarbonyl or C₂-Cβ alkylamino; C₃-Cβ cycloalkylamino; C₂-Cβ alkoxycarbonyl or C₂-Cβ alkylcarbonyl; or
- $R^2$  and  $R^3$  can be taken together with the nitrogen to which they are attached to form a ring containing 2 to 6 atoms of carbon and optionally one additional atom of nitrogen, sulfur or oxygen, said ring may be optionally substituted with 1 to 4 substituents selected from the group consisting of  $C_1$ - $C_2$  alkyl, halogen, CN, NO<sub>2</sub> and  $C_1$ - $C_2$  alkoxy;
- each R<sup>4</sup> is independently H, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>6</sub> haloalkyl, C<sub>2</sub>-C<sub>6</sub> haloalkenyl, C<sub>2</sub>-C<sub>6</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO<sub>2</sub>, hydroxy, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy,

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 $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$  haloalkylsulfinyl,  $C_1$ - $C_4$  haloalkylsulfinyl,  $C_1$ - $C_4$  haloalkylsulfonyl,  $C_1$ - $C_4$  alkylamino,  $C_2$ - $C_8$  dialkylamino,  $C_3$ - $C_6$  cycloalkylamino, or  $C_3$ - $C_6$  trialkylsilyl; or

- each R<sup>4</sup> is independently phenyl, benzyl or phenoxy, each optionally substituted with C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>2</sub>-C<sub>4</sub> haloalkenyl, C<sub>2</sub>-C<sub>4</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, C<sub>3</sub>-C<sub>6</sub> (alkyl)cycloalkylamino, C<sub>2</sub>-C<sub>4</sub> alkylcarbonyl, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub> dialkylaminocarbonyl or C<sub>3</sub>-C<sub>6</sub> trialkylsilyl;

(R<sup>5</sup>)<sub>2</sub> attached to adjacent carbon atoms can be taken together as OCF<sub>2</sub>O , CF<sub>2</sub>CF<sub>2</sub>O , or OCF<sub>2</sub>CF<sub>2</sub>O ;

- each R<sup>6</sup> is independently II, halogen, C<sub>1</sub>-C<sub>6</sub>-alkyl, C<sub>2</sub>-C<sub>6</sub>-alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>2</sub>-C<sub>6</sub> eyeloalkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy or C<sub>2</sub>-C<sub>4</sub> alkoxycarbonyl, or
- each R<sup>6</sup> is independently a phonyl, benzyl, phonoxy, 5 or 6 membered heteroaromatic ring or an aromatic 8, 9- or 10 membered fused heterobicyclic ring system, each ring optionally substituted with one to three substituents independently selected from the group consisting of C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> eyeloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>2</sub>-C<sub>4</sub> haloalkenyl, C<sub>2</sub>-C<sub>4</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> haloeyeloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>3</sub>-C<sub>6</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub> dialkylaminocarbonyl or C<sub>3</sub>-C<sub>6</sub> trialkylsilyl;
- each R<sup>7</sup> is independently H, C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>6</sub> haloalkyl, C<sub>2</sub>-C<sub>6</sub> haloalkenyl, C<sub>2</sub>-C<sub>6</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, CO<sub>2</sub>H, CONH<sub>2</sub>, NO<sub>2</sub>, hydroxy, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> haloalkylthio, C<sub>1</sub>-C<sub>4</sub> haloalkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> haloalkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>6</sub> dialkylamino, C<sub>2</sub>-C<sub>6</sub> alkylamino, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub> dialkylsilyl; or

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each R<sup>7</sup> is independently a phenyl, benzyl, benzoyl, phenoxy or 5- or 6-membered heteroaromatic ring or an 8-, 9- or 10-membered fused heterobicyclic ring system, each ring optionally substituted with one to three substituents independently selected from the group consisting of C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>2</sub>-C<sub>4</sub> haloalkenyl, C<sub>2</sub>-C<sub>4</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, C<sub>3</sub>-C<sub>6</sub> (alkyl)cycloalkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylaminocarbonyl, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub> dialkylaminocarbonyl and C<sub>3</sub>-C<sub>6</sub> trialkylsilyl;

## provided that

- (i) at least one R<sup>4</sup> and at least one R<sup>7</sup>, when R<sup>7</sup> is present, are other than H;
- (ii) J is other than an optionally substituted 1,2,3 thindiazole;
- (iii) (ii) when J is an optionally substituted pyridine and R<sup>2</sup> is H, R<sup>3</sup> is other than H or CH<sub>3</sub>;
- (iv) (iii) when J is an optionally substituted pyridine, then R<sup>7</sup> cannot be CONH<sub>2</sub>, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl or C<sub>3</sub>-C<sub>8</sub> dialkylaminocarbonyl; and
- (v) (iv) when J is an optionally substituted pyrazole, tetrazole or pyrimidine, then  $\mathbb{R}^2$  and  $\mathbb{R}^3$  cannot both be hydrogen; and
- (vi) when J is an optionally substituted 5 membered heterearomatic ring, then R<sup>2</sup> and R<sup>3</sup> are taken together with the nitrogen to which they are attached to form a ring containing 2 to 6 atoms of carbon and optionally one additional atom of nitrogen, sulfur or oxygen, said ring may be optionally substituted with 1 to 4 substituents selected from the group consisting of C<sub>1</sub> -C<sub>2</sub> alkyl, halogen, CN, NO<sub>2</sub> and C<sub>1</sub> -C<sub>2</sub> alkoxy.
  - 23. (Withdrawn): The compound of Claim 22 wherein J is a phenyl group substituted with 1 to 2 R<sup>5</sup> and optionally substituted with 1 to 3 R<sup>6</sup>.
  - 24. (Withdrawn): The compound of Claim 23 wherein

A and B are both O;

n is 1 to 2:

- $R^1$  is H,  $C_1$ - $C_4$  alkyl,  $C_2$ - $C_4$  alkenyl,  $C_2$ - $C_6$  alkynyl,  $C_3$ - $C_6$  cycloalkyl,  $C_2$ - $C_6$  alkylcarbonyl or  $C_2$ - $C_6$  alkoxycarbonyl;
- R<sup>2</sup> is H, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>2</sub>-C<sub>6</sub> alkylcarbonyl or C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl;
- R<sup>3</sup> is C<sub>1</sub>-C<sub>6</sub> alkyl, C<sub>2</sub>-C<sub>6</sub> alkenyl, C<sub>2</sub>-C<sub>6</sub> alkynyl or C<sub>3</sub>-C<sub>6</sub> cycloalkyl each optionally substituted with one or more substituents selected from the group consisting of halogen, CN, C<sub>1</sub>-C<sub>2</sub> alkoxy, C<sub>1</sub>-C<sub>2</sub> alkylthio, C<sub>1</sub>-C<sub>2</sub> alkylsulfinyl and C<sub>1</sub>-C<sub>2</sub> alkylsulfonyl;

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- one of the  $R^4$  groups is attached to the phenyl ring at the 2-position or 5-position, and said  $R^4$  is  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_4$  haloalkyl, halogen, CN, NO<sub>2</sub>,  $C_1$ - $C_4$  alkoxy,  $C_1$ - $C_4$  haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  haloalkylthio,  $C_1$ - $C_4$  haloalkylsulfinyl or  $C_1$ - $C_4$  haloalkylsulfonyl;
- each  $R^5$  is independently  $C_1$ - $C_4$  haloalkyl, CN, NO<sub>2</sub>,  $C_1$ - $C_4$  haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$  haloalkylsulfinyl,  $C_1$ - $C_4$  haloalkylsulfonyl or  $C_2$ - $C_4$  alkoxycarbonyl; or
- (R<sup>5</sup>)<sub>2</sub> when attached to adjacent carbon atoms can be taken together as -OCF<sub>2</sub>O-, -CF<sub>2</sub>CF<sub>2</sub>O- or -OCF<sub>2</sub>CF<sub>2</sub>O-; and
- each  $R^6$  is independently H, halogen,  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_2$  alkoxy or  $C_2$ - $C_4$  alkoxycarbonyl, or
- each R6 is independently a phenyl or a 5- or 6-membered heteroaromatic ring, each ring optionally substituted with C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>2</sub>-C<sub>4</sub> haloalkenyl, C<sub>2</sub>-C<sub>4</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylthio, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, C<sub>1</sub>-C<sub>4</sub> alkylamino, C<sub>2</sub>-C<sub>8</sub> dialkylamino, C<sub>3</sub>-C<sub>6</sub> cycloalkylamino, C<sub>3</sub>-C<sub>6</sub> (alkyl)cycloalkylamino, C<sub>2</sub>-C<sub>4</sub> alkylcarbonyl, C<sub>2</sub>-C<sub>6</sub> alkoxycarbonyl, C<sub>2</sub>-C<sub>6</sub> alkylaminocarbonyl, C<sub>3</sub>-C<sub>8</sub> dialkylaminocarbonyl or C<sub>3</sub>-C<sub>6</sub> trialkylsilyl.
- 25. (Withdrawn): The compound of Claim 24 wherein R<sup>1</sup> and R<sup>2</sup> are both H;
- R<sup>3</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with halogen, CN, OCH<sub>3</sub>, S(O)<sub>p</sub>CH<sub>3</sub>; each R<sup>4</sup> is independently H, CH<sub>3</sub>, CF<sub>3</sub>, OCF<sub>3</sub>, OCHF<sub>2</sub>, S(O)<sub>p</sub>CF<sub>3</sub>, S(O)<sub>p</sub>CHF<sub>2</sub>, CN or halogen;
- each R<sup>5</sup> is independently CF<sub>3</sub>, OCF<sub>3</sub>, OCHF<sub>2</sub>, S(O)<sub>p</sub>CF<sub>3</sub>, S(O)<sub>p</sub>CHF<sub>2</sub>, OCH<sub>2</sub>CF<sub>3</sub>, OCF<sub>2</sub>CHF<sub>2</sub>, S(O)<sub>p</sub>CH<sub>2</sub>CF<sub>3</sub> or S(O)<sub>p</sub>CF<sub>2</sub>CHF<sub>2</sub>;
- each R<sup>6</sup> is independently H, halogen or methyl; or phenyl, pyrazole, imidazole, triazole, pyridine or pyrimidine, each ring optionally substituted with C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, halogen or CN; and

p is 0, 1 or 2.

- 26. (Withdrawn): The compound of Claim 25 wherein R<sup>3</sup> is *i*-propyl or *t*-butyl.
- 27. (Canceled).
- 28. (Currently Amended): The compound of Claim 27 22 wherein

  J is a 5 or 6-membered heteroaromatic ring selected from the group consisting of J. 1, J. 2, J. 3, J. 4 and J. 5, J. 1 and J. 2 optionally substituted with 1 to 3

  R<sup>7</sup> and J. 3, wherein J. 4 and J. 5 substituted with R<sup>7</sup>

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Q is O, S or NR7; and

W, X, Y and Z are independently N or CR7, provided that in J 4 and J-5 at least one of W, X, Y or Z is N.

(Currently Amended): The compound of Claim 27 or Claim 28 wherein 29. A and B are O;

n is 1 to 2:

 $R^1$  is H,  $C_1$ - $C_4$  alkyl,  $C_2$ - $C_4$  alkenyl,  $C_2$ - $C_4$  alkynyl,  $C_2$ - $C_6$  alkylcarbonyl or  $C_2$ - $C_6$ alkoxycarbonyl;

R<sup>2</sup> is H, C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>2</sub>-C<sub>4</sub> alkenyl, C<sub>2</sub>-C<sub>4</sub> alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>2</sub>-C<sub>6</sub> alkylcarbonyl or C2-C6 alkoxycarbonyl;

R3 is H; or C1-C6 alkyl, C2-C6 alkenyl, C2-C6 alkynyl or C3-C6 cycloalkyl each optionally substituted with one or more substituents selected from the group consisting of halogen, CN, C1-C2 alkoxy, C1-C2 alkylthio, C1-C2 alkylsulfinyl and C1-C2 alkylsulfonyl;

one of the R4 groups is attached to the phenyl ring at the 2-position, and said R4 is C1-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, halogen, CN, NO<sub>2</sub>, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$  haloalkylthio,  $C_1$ - $C_4$  haloalkylsulfinyl or  $C_1$ - $C_4$  haloalkylsulfonyl; and

each  $\mathbb{R}^7$  is independently H,  $\mathbb{C}_1$ - $\mathbb{C}_4$  alkyl,  $\mathbb{C}_1$ - $\mathbb{C}_4$  haloalkyl, halogen, CN,  $\mathbb{NO}_2$ ,  $\mathbb{C}_1$ - $\mathbb{C}_4$ haloalkoxy,  $C_1$ - $C_4$  alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$ haloalkylthio,  $C_1$ - $C_4$  haloalkylsulfinyl,  $C_1$ - $C_4$  haloalkylsulfonyl or C2-C4 alkoxycarbonyl; or a phenyl or a 5- or 6-membered heteroaromatic ring, each ring optionally substituted with C1-C4 alkyl, C2-C4 alkenyl, C2-C4 alkynyl, C<sub>3</sub>-C<sub>6</sub> cycloalkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>2</sub>-C<sub>4</sub> haloalkenyl, C<sub>2</sub>-C<sub>4</sub> haloalkynyl, C<sub>3</sub>-C<sub>6</sub> halocycloalkyl, halogen, CN, NO2,  $C_1$ - $C_4$  alkoxy,  $C_1$ - $C_4$  haloalkoxy,  $C_1$ - $C_4$ alkylthio,  $C_1$ - $C_4$  alkylsulfinyl,  $C_1$ - $C_4$  alkylsulfonyl,  $C_1$ - $C_4$  alkylamino,  $C_2$ - $C_8$ dialkylamino, C3-C6 cycloalkylamino, C3-C6 (alkyl)cycloalkylamino, C2-C4 alkylearbonyl, C2-C6 alkoxycarbonyl, C2-C6 alkylaminocarbonyl, C3-C8 dialkylaminocarbonyl or  $C_3$ - $C_6$  trialkylsilyl.

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- 30. (Currently Amended): The compound of Claim 29 wherein J is selected from the group consisting of pyridine, and pyrimidine, pyrazole, imidazole, triazoles, thiophene and thiazole, each optionally substituted with 1 to 3 R<sup>7</sup>.
- 31. (Currently Amended): The compound of Claim 30 wherein
- J is selected from the group consisting of pyridine, and pyrimidine, pyrazole, thiophene and thiazole, each optionally substituted with 1 to 3 R<sup>7</sup>;

R1 and R2 are both H;

R<sup>3</sup> is C<sub>1</sub>-C<sub>4</sub> alkyl optionally substituted with halogen, CN, OCH<sub>3</sub> or S(O)<sub>p</sub>CH<sub>3</sub>; each R<sup>4</sup> is independently H, CH<sub>3</sub>, CF<sub>3</sub>, OCF<sub>3</sub>, OCHF<sub>2</sub>, S(O)<sub>p</sub>CF<sub>3</sub>, S(O)<sub>p</sub>CHF<sub>2</sub>, CN or halogen;

each R7 is independently H, halogen, CH<sub>3</sub>, CF<sub>3</sub>, OCHF<sub>2</sub>, S(O)<sub>p</sub>CF<sub>3</sub>, S(O)<sub>p</sub>CHF<sub>2</sub>, OCH<sub>2</sub>CF<sub>3</sub>, OCF<sub>2</sub>CHF<sub>2</sub>, S(O)<sub>p</sub>CH<sub>2</sub>CF<sub>3</sub>, or S(O)<sub>p</sub>CF<sub>2</sub>CHF<sub>2</sub>; or phenyl, pyrazole, imidazole, triazole, pyridine or pyrimidine, each ring optionally substituted with C<sub>1</sub>-C<sub>4</sub> alkyl, C<sub>1</sub>-C<sub>4</sub> haloalkyl, C<sub>1</sub>-C<sub>4</sub> alkoxy, C<sub>1</sub>-C<sub>4</sub> haloalkoxy, C<sub>1</sub>-C<sub>4</sub> alkylsulfinyl, C<sub>1</sub>-C<sub>4</sub> alkylsulfonyl, halogen or CN; and p is 0, 1 or 2.

- 32. (Original): The compound of Claim 31 wherein J is a pyridine optionally substituted with 1 to 3 R<sup>7</sup>.
- 33. (Original): The compound of Claim 32 wherein one  $\mathbb{R}^7$  is a phenyl optionally substituted with  $\mathbb{C}_1$ - $\mathbb{C}_4$  alkyl,  $\mathbb{C}_1$ - $\mathbb{C}_4$  haloalkyl, halogen or  $\mathbb{C}\mathbb{N}$ .
- 34. (Original): The compound of Claim 32 wherein one  $R^7$  is a pyrazole, imidazole, triazole, pyridine or pyrimidine, each ring optionally substituted with  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_4$  haloalkyl, halogen or CN.
- 35. (Original): The compound of Claim 31 wherein J is a pyrimidine optionally substituted with 1 to 3 R<sup>7</sup>.
- 36. (Original): The compound of Claim 35 wherein one  $\mathbb{R}^7$  is a phenyl optionally substituted with  $\mathbb{C}_1$ - $\mathbb{C}_4$  alkyl,  $\mathbb{C}_1$ - $\mathbb{C}_4$  haloalkyl, halogen or  $\mathbb{C}\mathbb{N}$ .
- 37. (Original): The compound of Claim 35 wherein one  $R^7$  is a pyrazole, imidazole, triazole, pyridine or pyrimidine, each ring optionally substituted with  $C_1$ - $C_4$  alkyl,  $C_1$ - $C_4$  haloalkyl, halogen or CN.
  - 38. (Canceled).
  - 39. (Canceled).
  - 40. (Canceled).
  - 41. (Canceled).
- 42. (Currently Amended): The compound of Claim 22 selected from the group consisting of: which is

3 methyl-N-(1-methylethyl)-2 [[4-(trifluoromethyl)benzoyl]amino]-benzamide,

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2-methyl-N [2-methyl-6 [[(1-methylethyl)mnino]carbonyl]phenyl]-4-(trifluoromethyl)benzamide, <u>und</u> 2-methyl-N-[2-methyl-6-[[(1-methylethyl)amino]carbonyl]phenyl]-6-(trifluoromethyl)-3-pyridinecarboxamide.

43. (Original): An arthropodicidal composition comprising an arthropodicidally effective amount of a compound of Formula 1 as described in Claim 1 and at least one additional component selected from the group consisting of surfactants, solid diluents and liquid diluents.